



DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Health Service

Food and Drug Administration Southwest Region 7920 Elmbrook Drive Suite 102 Dallas, TX 75247-4982

Telephone: 214-655-8100 FAX: 214-655-8130

January 16, 2002

WARNING LETTER

CERTIFIED MAIL RETURN RECEIPT REQUESTED

02-SWR-WL-23/7

Bob McCormack Technical Director of Radiology North Hills Hospital Subsidiary, L.P., Inc. dba North Hills Hospital 4401 Booth Calloway Road North Richland Hills, TX 76180

RE: Inspection ID - 1276390007

Dear Mr. McCormack,

On 1/9/2002, a representative from the state of Texas, acting on behalf of the Food and Drug Administration (FDA) inspected your facility. This inspection revealed a serious regulatory problem involving the mammography at your facility.

The Mammography Quality Standards Act of 1992 requires your facility to meet specific standards. These requirements help protect the health of women by assuring that a facility can perform quality mammography. The inspection revealed the following level 1 finding at your facility:

- Level 1: The system to communicate results is not adequate for site North Hills Hospital Subsidiary, L.P., Inc. because:
- There is no system in place to provide timely lay summaries

The specific problem noted above appeared on your MQSA Facility Inspection Report, which was issued to your facility at the close of the inspection.

Level 1 findings may be symptomatic of serious underlying problems that could compromise the quality of mammography at your facility. They represent a serious violation of the law which may result in FDA taking regulatory action without further notice to you. These actions include, but are not limited to:

- Placing your facility under a Directed Plan of Correction.
- Charging your facility for the cost of on-site monitoring.

- Assessing civil money penalties up to \$10,000 for each failure to substantially comply with, or each day of failure to substantially comply with, the Standards.
- Suspension or revocation of your facility's FDA certificate, or obtaining a court injunction against further mammography.

In addition, your response should address the Level 2 findings that were listed on the inspection report provided to you at the close of the inspection. These Level 2 findings are:

- Level 2: Medical audit and outcome analysis was not done separately for each individual at site North Hills Hospital Subsidiary, L.P., Inc.

It is necessary for you to act on this matter immediately. You are required to respond to this office in writing within fifteen (15) working days from receipt of this letter. Please address the following:

- The specific steps you have taken to correct all of the violations noted in this letter.
- Each step your facility is taking to prevent the recurrence of similar violations.
- Equipment settings (including technique factors), raw test data, and calculated final results, where appropriate.
- Sample records that demonstrate proper record keeping procedures, if the findings relate to quality control or other records (Note: Patient names or identification should be deleted from any copies submitted).

Please submit your response to: Deborah McGee, Radiation Specialist Food and Drug Administration 7920 Elmbrook Drive, Suite 102 Dallas, Texas 75247-4982

This letter pertains only to findings of your inspection and does not necessarily address other obligations you have under the law. You may obtain general information about all of FDA's requirements for mammography facilities by contacting the Mammography Quality Assurance Program, Food and Drug Administration, P.O. Box 6057, Columbia, MD 21045-6057 (1-800-838-7715) or through the Internet at http://www.fda.gov/cdrh/mammography/index.html.

If you have more specific questions about mammography facility requirements, or about the content of this letter, please feel free to contact Angela Moak at (214) 655-8100 ext. 135.

Sincerely yours,

Gary L. Pierce
Regional Food and Drug Director